Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

 (Previously presented) A method of identifying an agent that binds to CCX-CKR2 on a cell, the method comprising,

contacting a plurality of agents and a CCX-CKR2 ligand to a CCX-CKR2 polypeptide comprising an extracellular domain at least 95% identical to an extracellular domain of a CCX-CKR2 polypeptide comprising SEQ ID NO:2, or a fragment of the CCX-CKR2 polypeptide that binds SDF1 or I-TAC, wherein the CCX-CKR2 ligand is SDF1 or I-TAC; and

selecting an agent that competes with I-TAC or SDF1 for binding to the CCX-CKR2 polypeptide or fragment thereof, thereby identifying an agent that binds to CCX-CKR2 on a cell.

- 2. (Original) The method of claim 1, wherein the cell is a cancer cell.
- 3. (Original) The method of claim 1, further comprising testing the selected agent for the ability to bind to, or inhibit growth of, a cell.
 - 4. (Original) The method of claim 3, wherein the cell is a cancer cell.
- (Original) The method of claim 1, further comprising testing the selected agent for the ability to alter kidney function.
- (Original) The method of claim 1, further comprising testing the selected agent for the ability to alter brain or neuronal function.
- (Original) The method of claim 1, further comprising testing the selected agent for the ability to change cell adhesion to endothelial cells.
- 8. (Original) The method of claim 1, wherein the agent is less than 1,500 daltons.

PATENT

Appl. No. 10/698,541 Amdt. dated March 6, 2007 Reply to Office Action of February 1, 2007

- 9. (Original) The method of claim 1, wherein the agent is an antibody.
- (Original) The method of claim 1, wherein the CCX-CKR2 polypeptide comprises the sequence displayed in SEQ ID NO:2.
 - 11-27. (Canceled)
- (Previously presented) A method of competing SDF1 or I-TAC and an agent for binding to a CCX-CKR2 polypeptide, the method comprising

contacting a cell with an agent that specifically binds to a polypeptide comprising SEQ ID NO:2, wherein the agent competes with SDF-1 or I-TAC for binding to the CCX-CKR2 polypeptide, and wherein the cell expresses a CCX-CKR2 polypeptide comprising an extracellular domain at least 95% identical to an extracellular domain of SEQ ID NO:2.

- (Original) The method of claim 28, wherein the agent is less than 1,500 daltons.
 - 30. (Original) The method of claim 28, wherein the agent is an antibody.
- (Original) The method of claim 28, wherein the CCX-CKR2 polypeptide is as displayed in SEQ ID NO:2.
- (Original) The method of claim 28, wherein the agent is identified by a
 method comprising

contacting a plurality of agents to a CCX-CKR2 polypeptide comprising an extracellular domain at least 95% identical to an extracellular domain of a CCX-CKR2 polypeptide comprising SEQ ID NO:2, or a fragment of the CCX-CKR2 polypeptide that binds SDF1 or I-TAC: and

selecting an agent that competes with I-TAC or SDF-1 for binding to the CCX-CKR2 polypeptide or fragment thereof, thereby identifying an agent that binds to a cancer cell.

33-38. (Canceled)

(Currently amended) The method of claim 1 2, wherein the CCX-CKR2
 ligand is detectably-labeled and the selecting step comprises measuring the amount of labeled

Appl. No. 10/698,541

Amdt. dated March 6, 2007

Reply to Office Action of February 1, 2007

CCX-CKR2 ligand bound to the polypeptide in the presence of at least one of the plurality of agents.